Exhibit 5 510(k) Summary

AUG 26 2011

Picture archiving and communications system / Model: Zenis

1. Company and Correspondent making the submission

1.1 Submitter and US Official Correspondent

Submitter: GENORAY Co., Ltd.
Address: #512, Byucksan Technopia, 434-6, Sangdaewon 1-dong, Jungwon-gu, Seongnam-city, Gyeonggi-do, 462-716 Korea
Telephone No.: +82-31-740-4100
Fax: +82-31-737-8025

1.2 Official Correspondent (U.S): Jae Kim - Business Manager

Correspondent: GENORAY America Inc.
Address: 1073 N. Batavia St. Orange, CA 92867, USA
Telephone No.: 714-289-8020
Fax: 714-453-9661
Email: jae@genoray.com

2. Establishment Registration Number

3005843418

3. Device Information

Proprietary/Trade Name:	Picture archiving and communications system / Model: Zenis
Common/Usual Name:	Picture archiving and communications system
Classification Name:	System, Image Processing, Radiological
Product Code:	LLZ
Device Class:	Class II per regulation 21 CFR 892.2050

4. Equivalent Legally Marketed Device

Manufacturer:	GE Medical Systems Information Technologies
Device Name:	RA600
510(k) Number:	K042525 (Decision Date – October 1, 2004)
Classification:	System, Image Processing, Radiological: LLZ, Class II per regulation 21 CFR 892.2050

5. Description of the Device

Zenis is a PC-based DICOM workstation platform which provides scalable image and data management solutions for medical imaging. This software-based product provides capabilities for the acceptance, transmission, printing, display, storage, editing and digital processing of medical images and associated data.

Zenis may be combined with a PACS network or connected directly to a modality through the use of DICOM networking.

6. Indications for use

Zenis is intended for viewing of images acquired from Fluoroscopic X-ray system when installed on suitable commercial-standard PC hardware. Zenis is intended for use as a primary diagnostic and analysis workstation in Radiology or other departments. It is also intended for use as a clinical review workstation throughout the healthcare facility and may be part of a larger PACS configuration.

7. Safety and Effectiveness, comparison to Predicate

Zenis, which is made by GENORAY Co., Ltd., is substantially equivalent to RA600 of GE healthcare. We selected the RA600 as the predicate device already FDA approved. Because of RA600 and Zenis are almost same in function & characteristic. Zenis & RA600 are intended for use as PACS software used for viewing of medical images acquired from modality when installed on suitable commercial-standard PC hardware.

The result of performance test and clinical evaluation indicates that the new device is as safe and effective as the predicate devices.

8. Conclusion

In accordance with the Federal Food, Drug and Cosmetic Act, 21 CFR Part 807 GENORAY Co., Ltd., concludes that the Picture archiving and communications system (Model: Zenis) is safe and effective and substantially equivalent to the predicate device as described above.

DEPARTMENT OF HEALTH & HUMAN SERVICES Public Health Service

Food and Drug Administration
10903 New Hampshire Avenue
Document Control Room – WO66-G609
Silver Spring, MD 20993-0002

Genoray Co., Ltd.
% Mr. Jae Kim
Business Development Manager
Genoray America, Inc.
1073 N. Batavia St.
ORANGE CA 92867

AUG 26 2011

Re: K103181
Trade/Device Name: Picture archiving and communications system (Models: Zenis)
Regulation Number: 21 CFR 892.2050
Regulation Name: Picture archiving and communications system
Regulatory Class: II
Product Code: LLZ
Dated: October 15, 2010
Received: July 8, 2011

Dear Mr. Kim:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into class II (Special Controls), it may be subject to such additional controls. Existing major regulations affecting your device can be found in Title 21, Code of Federal Regulations (CFR), Parts 800 to 895. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Parts 801 and 809); medical device reporting (reporting of

medical device-related adverse events) (21 CFR 803); and good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820). This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Parts 801 and 809), please contact the Office of *In Vitro* Diagnostic Device Evaluation and Safety at (301) 796-5450. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely Yours,

Mary S Pastel

Mary S. Pastel, Sc.D.
Director
Division of Radiological Devices
Office of In Vitro Diagnostic Device
Evaluation and Safety
Center for Devices and Radiological Health

Enclosure

Exhibit 4 Indications for use

510(k) number (if known): K103181

Device Name: Picture archiving and communications system (Models: Zenis)

Indications for Use:

Zenis is intended for viewing of images acquired from Fluoroscopic X-ray system when installed on suitable commercial-standard PC hardware. Zenis is intended for use as a primary diagnostic and analysis workstation in Radiology or other departments. It is also intended for use as a clinical review workstation throughout the healthcare facility and may be part of a larger PACS configuration.

Prescription Use ___V___ AND/OR Over-The-Counter Use _____

(Part 21 CFR 801 Subpart D) (Part 21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Radiological Devices
Office of In Vitro Diagnostic Device Evaluation and Safety

510K K103181